

EMETROL CHERRY- phosphorated carbohydrate solution
WellSpring Pharmaceutical Corporation

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).

Emetrol

Active ingredients (per 5mL)

Dextrose (glucose) 1.87g

Levulose (fructose) 1.87g

Phosphoric acid 21.5mg

Purpose

Upset Stomach Reliever

Uses

- for relief of upset stomach associated with nausea

Warnings

- This product contains fructose and should not be taken by persons with hereditary fructose intolerance (HFI).

Do not use if you have

- allergic reactions to any of the ingredients in this product

Ask a doctor before use if you have

- diabetes

Stop use and ask a doctor if

- symptoms persist, return or get worse

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

- for maximum effectiveness never dilute or drink fluids of any kind immediately before or after taking this product

- repeat dose every 15 minutes or until distress subsides
- do not take more than 5 doses in 1 hour without consulting a doctor
- measure only with dosing cup provided. Dosing cup to be used with product only. Do not use with other products.
- mL = milliliters

Age	Dose
Adults and children 12 years of age and over	15 mL or 30 mL
Children 2 to under 12 years of age	5 mL or 10 mL

Other information

- Store between 15-30°C (59-86°F) away from heat and direct light; keep from freezing
- **Do not use if printed foil seal under bottle cap is broken or missing**

Inactive ingredients

FD&C red no. 40, flavors, glycerin, methylparaben, and purified water.

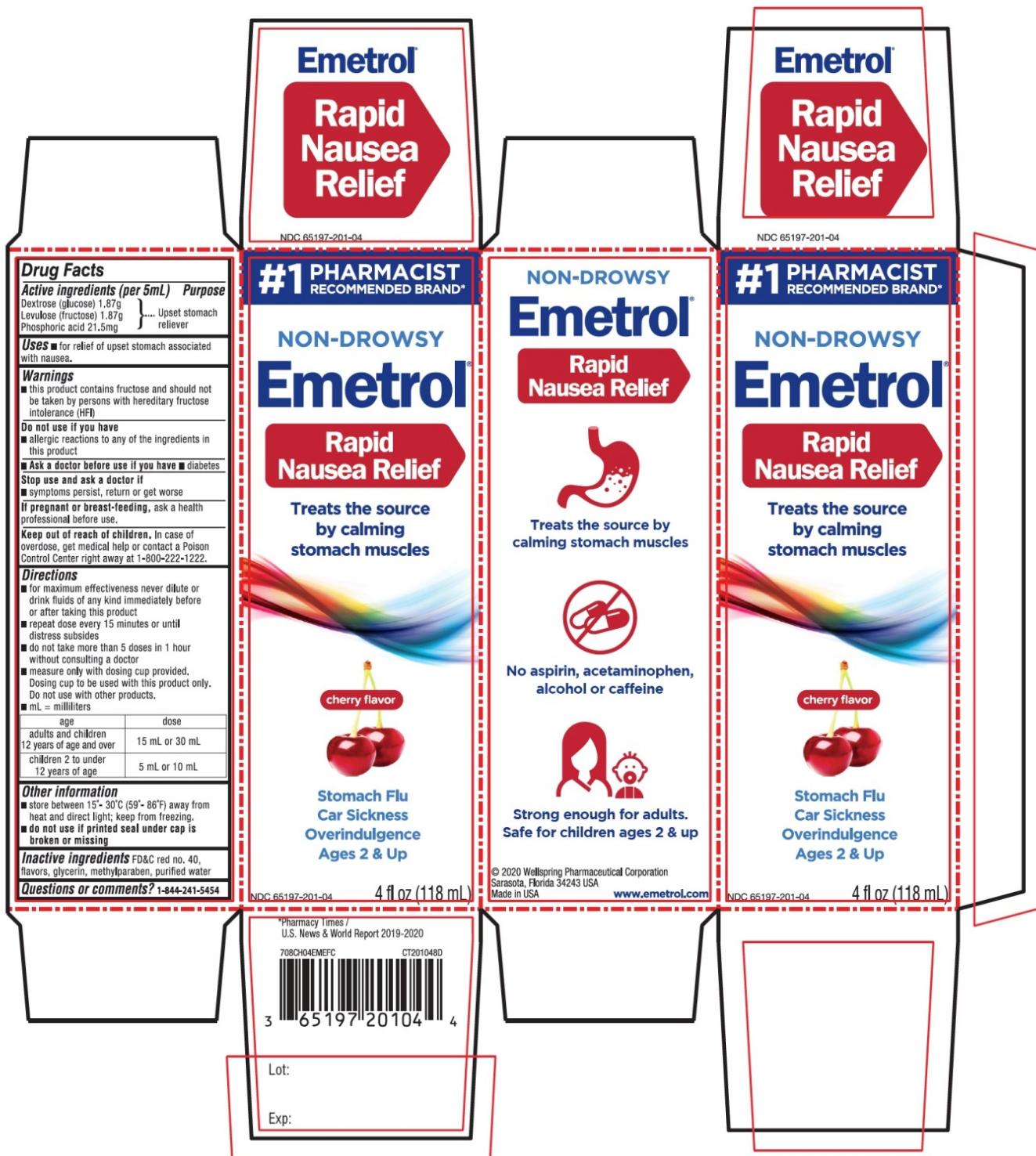
Questions or Comments?

call 1-844-241-5454

Distributed By

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PACKAGE LABEL



EMETROL CHERRY

phosphorated carbohydrate solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65197-201
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2) (DEXTROSE, UNSPECIFIED FORM - UNII:IY9XDZ35W2)			DEXTROSE, UNSPECIFIED FORM	1.87 g in 5 mL
FRUCTOSE (UNII: 6YSS42VSEV) (FRUCTOSE - UNII:6YSS42VSEV)			FRUCTOSE	1.87 g in 5 mL
PHOSPHORIC ACID (UNII: E4GA8884NN) (PHOSPHORIC ACID - UNII:E4GA8884NN)			PHOSPHORIC ACID	21.5 mg in 5 mL
Inactive Ingredients				
Ingredient Name			Strength	
FD&C RED NO. 40 (UNII: WZB9127XOA)				
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
Product Characteristics				
Color	RED (Red)		Score	
Shape			Size	
Flavor	CHERRY		Imprint Code	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65197-201-04	1 in 1 CARTON	06/24/2009	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:65197-201-06	133 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2018	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other			06/24/2009	

Labeler - WellSpring Pharmaceutical Corporation (110999054)

Revised: 8/2020

WellSpring Pharmaceutical Corporation